

**AMENDMENT UNDER 37 C.F.R. § 1.116**  
**U.S. Appl. No. 10/533,833 (Q101061)**

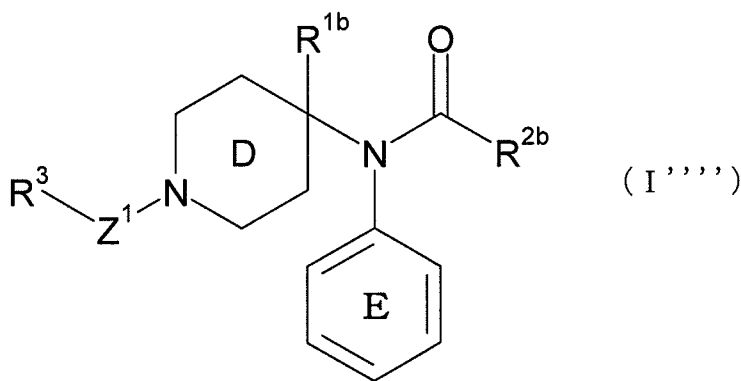
**AMENDMENTS TO THE CLAIMS**

**This listing of claims will replace all prior versions and listings of claims in the application:**

**LISTING OF CLAIMS:**

1-10. (cancelled).

11. (currently amended) A compound represented by the formula:



wherein ring D represents a piperidine ring ~~optionally further substituted with C<sub>1-6</sub> alkyl~~, E represents a phenyl group optionally substituted with a substituent selected from the group consisting of a halogen atom and C<sub>1-6</sub> alkyl, Z<sup>1</sup> represents a methylene group optionally substituted with ~~a substituent selected from the group consisting of C<sub>1-6</sub> alkyl, C<sub>1-6</sub> alkoxy, carbonyl, oxo and phenyl, COCH<sub>2</sub>, CH<sub>2</sub>CO or SO<sub>2</sub>~~, R<sup>1b</sup> represents (i) a 2-thiazolyl group optionally substituted with C<sub>1-6</sub> alkyl, (ii) a 2-imidazolyl group optionally substituted with C<sub>1-6</sub> alkyl, (iii) a 2-pyridyl group optionally substituted with a substituent selected from the group consisting of C<sub>1-6</sub> alkyl, a halogen atom, C<sub>1-6</sub> alkylthio, ~~phenyl~~ and thienyl, R<sup>2b</sup> represents an optionally halogenated C<sub>1-6</sub> alkyl group, and R<sup>3</sup> represents ~~(i) a C<sub>3-8</sub> cycloalkyl group, (ii) a phenyl group or (iii) a 5- to 10-membered aromatic heterocyclic group containing one or two~~

**AMENDMENT UNDER 37 C.F.R. § 1.116**  
**U.S. Appl. No. 10/533,833 (Q101061)**

~~kinds of 1 to 4 hetero atoms selected from a nitrogen atom, a sulfur atom and an oxygen atom in addition to carbon atoms, which may be substituted with a substituent selected from the group consisting of a halogen atom, cyano, C<sub>1-6</sub> alkyl optionally substituted with a halogen atom, C<sub>1-6</sub> alkoxy optionally substituted with a halogen atom, C<sub>1-6</sub> alkyl carbonylamino, a 5- or 6-membered aromatic heterocyclic group and C<sub>1-6</sub> alkylthio, provided that 1) N-[1-benzyl-4-(thiazol-2-yl)-4-piperidinyl]-N-phenylpropionamide, 2) N-[1-benzyl-4-(thiazol-2-yl)-4-piperidinyl]-N-(2-fluorophenyl)propionamide, 3) N-[1-benzyl-4-(4-methylthiazol-2-yl)-4-piperidinyl]-N-(2-fluorophenyl)propionamide, 4) N-[1-benzyl-4-(4,5-dimethylthiazol-2-yl)-4-piperidinyl]-N-phenylpropionamide, 5) N-[1-benzyl-4-(4,5-dimethylthiazol-2-yl)-4-piperidinyl]-N-(2-fluorophenyl)propionamide, 6) N-[1-benzyl-4-(2-pyridinyl)-4-piperidinyl]-N-(2-fluorophenyl)propionamide, 7) N-[1-benzyl-4-(4-methylthiazol-2-yl)-4-piperidinyl]-N-phenylpropionamide and 8) N-[1-benzyl-4-(2-pyridinyl)-4-piperidinyl]-N-phenylpropionamide are excluded, or a salt thereof.~~

12. (cancelled).

13. (original) The compound according to claim 11, wherein R<sup>3</sup> is a phenyl group.

14. (original) The compound according to claim 11, wherein E is a phenyl group optionally having a substituent at an ortho position or a meta position.

15. (original) The compound according to claim 11, wherein E is an unsubstituted phenyl group.

**AMENDMENT UNDER 37 C.F.R. § 1.116**  
**U.S. Appl. No. 10/533,833 (Q101061)**

16. (previously presented) The compound according to claim 11, wherein R<sup>1b</sup> is a 2-thiazolyl group optionally substituted with a C<sub>1-6</sub> alkyl group.

17. (original) The compound according to claim 11, wherein R<sup>1b</sup> is a 4-methyl-2-thiazolyl group.

18. (withdrawn-previously presented) The compound according to claim 11, wherein R<sup>1b</sup> is a 2-pyridyl group optionally substituted with a substituent selected from the group consisting of a C<sub>1-6</sub> alkyl group, a C<sub>1-6</sub> alkylthio group, a halogen atom, a C<sub>6-14</sub> aryl group and an aromatic heterocyclic group.

19. (withdrawn) The compound according to claim 11, wherein R<sup>1b</sup> is a 6-methyl-2-pyridyl group.

20. (cancelled).

21. (original) The compound according to claim 11, wherein Z<sup>1</sup> is a methylene group.

22. (original) The compound according to claim 11, wherein R<sup>2b</sup> is an optionally halogenated methyl group or ethyl group.

23. (original) The compound according to claim 11, wherein R<sup>2b</sup> is a methyl group or a trifluoromethyl group.

**AMENDMENT UNDER 37 C.F.R. § 1.116**  
**U.S. Appl. No. 10/533,833 (Q101061)**

24-25. (cancelled).

26. (previously presented) N-[1-benzyl-4-(4-methylthiazol-2-yl)-4-piperidinyl]-N-phenylacetamide, N-[1-benzyl-4-(4-methylthiazol-2-yl)-4-piperidinyl]-2,2,2-trifluoro-N-phenylacetamide, N-[1-benzyl-4-(6-methyl-2-pyridinyl)-4-piperidinyl]-N-phenylacetamide, N-[1-benzyl-4-(6-methyl-2-pyridinyl)-4-piperidinyl]-2,2,2-trifluoro-N-phenylacetamide, N-[1-(4-fluorobenzyl)-4-(4-methylthiazol-2-yl)-4-piperidinyl]-N-phenylacetamide, N-[1-benzyl-4-(4-methylthiazol-2-yl)-4-piperidinyl]-N-(2-methylphenyl)acetamide, N-[1-benzyl-4-(4-methylthiazol-2-yl)-4-piperidinyl]-N-(3-chlorophenyl)acetamide, N-[4-(4-methylthiazol-2-yl)-1-(2-thienylmethyl)-4-piperidinyl]-N-phenylacetamide, N-[1-benzyl-4-(1-methyl-1H-imidazol-2-yl)-4-piperidinyl]-N-phenylacetamide, or a salt thereof.

27. (cancelled).

28. (previously presented) A medicine comprising the compound according to claim 11 or 26 or a salt thereof.

29. (previously presented) A pharmaceutical composition for regulating neuromedin U receptor, which comprises the compound according to claim 11 or 26 or a salt thereof and a pharmaceutically acceptable carrier.